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OM protein - protein search, using sw model

Run on: May 29, 2003, 15:21:44 ; Search time 22 Seconds

(Without alignments)
152.708 Million cell updates/sec

Title: US-09-924-102-2

Perfect score: 81
Sequence: 1 MLSTHFLYFLYFLYSYL.....RMGGGGRGGGTADTGWFLS 81

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 112892 seqs, 41476328 residues

Word size: 0

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database: SwissProt_40:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match Length	ID	Description
1	8	9.9	65	1 RPR_DROME	Q24475 drosophila
2	8	9.9	284	1 SUHA_HUMAN	Q06520 homo sapien
3	8	9.9	284	1 SUHA_MACFA	P52842 macaca fasc
4	8	9.9	587	1 ALU2_HUMAN	P39189 homo sapien
5	8	9.9	587	1 ALU3_HUMAN	P39190 homo sapien
6	8	9.9	591	1 ALU8_HUMAN	P39195 homo sapien
7	8	9.9	591	1 ALU8_HUMAN	P39195 homo sapien
8	8	9.9	116	1 FLHD_XENNE	Q39912 xenorhabdus
9	8	9.9	116	1 FLHD_XENNE	Q39912 xenorhabdus
10	8	9.9	285	1 INVO_CANFA	P18174 canis famli
11	8	9.9	304	1 Y687_HABIN	P11356 haemophilus
12	8	9.9	337	1 Y687_HABIN	P11356 haemophilus
13	8	9.9	352	1 CITC_ECOLI	P73164 saccharomyc
14	8	9.9	352	1 CITC_ECOLI	P73164 saccharomyc
15	8	9.9	366	1 ROX1_YEAST	P25042 saccharomyc
16	8	9.9	591	1 ALU1_HUMAN	P39188 homo sapien
17	8	9.9	603	1 ALU4_HUMAN	P39191 homo sapien
18	8	9.9	804	1 SYL_BACSU	P36430 bacillus su
19	8	9.9	806	1 SYL_BACSU	Q9K768 bacillus ha
20	8	9.9	889	1 C122_HUMAN	Q93826 homo sapien
21	8	9.9	1230	1 UGSA_SOLTU	Q43846 solanum tub
22	8	9.9	66	1 HSP1_ISOMA	P42136 isodon mac
23	8	9.9	68	1 HSP1_PERGU	P42137 perameles g
24	8	9.9	93	1 H1S2_MYCTU	Q33257 mycobacteri
25	8	9.9	118	1 RS20_SCHPO	Q07483 schistosach
26	8	9.9	134	1 SY21_HUMAN	Q00585 homo sapien
27	8	9.9	140	1 RL23_DROME	P48159 drosophila
28	8	9.9	153	1 IF1A_YEAST	P38912 drosophila
29	8	9.9	154	1 YK01_AERPE	Q9ya48 saccharomyc
30	8	9.9	160	1 HBL1_ARATH	Q24520 arabidopsis
31	8	9.9	168	1 RR7_CHLRE	P48267 chlamydomon
32	8	9.9	174	1 IL1X_BOVIN	Q77482 bos taurus
33	8	9.9	176	1 DPSA_SYMP7	Q55024 synecococc
34	8	9.9	189	1 COAT_TYMP	P03608 turnip yell
35	8	9.9	189	1 COAT_TYMP	P20125 turnip yell
36	8	9.9	193	1 KITH_HAEMIN	P44309 haemophilus

34	6	7.4	201	1 COAE_BACHD	Q9K857 bacillus ha
35	6	7.4	206	1 COX3_BACFI	Q04442 bacillus fi
36	6	7.4	211	1 CLUD_MOUSE	Q92064 mus musculu
37	6	7.4	214	1 GRP2_NICSY	P27484 nicotiana s
38	6	7.4	231	1 TRMD_MYCE	P47683 mycoplasma
39	6	7.4	252	1 TH14_PYRAB	Q9V018 pyrococcus
40	6	7.4	255	1 TH14_PYRHO	Q59082 pyrococcus
41	6	7.4	262	1 TRPA_AQUAE	Q67502 aquifex aeo
42	6	7.4	281	1 RP32_HABIN	P44404 haemophilus
43	6	7.4	284	1 RP32_CITFR	P11538 citrobacter
44	6	7.4	284	1 RP32_ECOLI	P00580 escherichia
45	6	7.4	285	1 RP32_ENTCL	P50508 enterobacte

ALIGNMENTS

RESULT 1
RPR_DROME STANDARD: PRT: 65 AA.
AC Q24475: Q9VAP7;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Reaper protein.
GN RPR OR CG4319.
OS Drosophila melanogaster (fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;
OC Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
OC Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_Taxid=7227;
[1]
RN
RP SEQUENCE FROM N.A.
RC STRAIN=Oregon-R; TISSUE=Eye Imaginal disk;
RX MEDLINE=94225205; PubMed=8171319;
RA White K., Grether M.E., Abrams J.M., Young L., Farrell K., Steller H.;
RT "Genetic control of programmed cell death in Drosophila.";
RL Science 264:677-683(1994).
[2]
RP SEQUENCE FROM N.A.
RC STRAIN=Berkley;
RX MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celisner S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Amandides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Wortman J.R., Lander M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazey R.G., Champe M., Pfeiffer B.D.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
RA Abril J.F., Aghayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Bernan B.P., Bhandari D., Bolshakov S.,
RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotter P.,
RA Butris K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davaport L.B., Davies P.,
RA De Pablo S., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Dou P.L.E., Downes M., Dugan-Rocha S., Dunkov B., Dunn P.,
RA Durbin K.J., Evangelista C.C., Ferraz C., Ferriere S., Fleischmann W.,
RA Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodex A., Gong F., Gottrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hottis D., Houston K.A., Howland T.J., Wei M.-H., Idegawa C.,
RA Jaitai M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Mishina N.V., Moberly B., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacle J.M.,
RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheefer F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Slier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,

RA Wang Z.-Y., Maasman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodard T., Morley R.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbe R.A., Myers E.W., Rubin G.M., Venter J.C.,
 RT "The genome sequence of Drosophila melanogaster.",
 RL Science 287:2185-2195(2000).
 CC -1- FUNCTION: PLAYS A CENTRAL AND GLOBAL REGULATORY FUNCTION FOR THE
 CC INITIATION OF APOPTOSIS. ECTOPIC EXPRESSION IN THE DEVELOPING EYE
 CC RESULTS IN A SMALL EYE DUE TO EXCESS CELL DEATH.
 CC -1- DEVELOPMENTAL STAGE: EXPRESSION CORRESPONDS TO THE PATTERN OF
 CC PROGRAMMED CELL DEATH IN THE EMBryo.
 CC -1- SIMILARITY: LIMITED AT THE N-TERMINAL, TO HID AND GRIM.
 CC
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 CC
 DR EMBL: L31631; AAA18983.1; -
 DR EMBL: AE003520; AAF49264.1; -
 DR FlyBase: FBgn0011706; rpr.
 DR Apoptosis.
 KW SEQUENCE 65 AA; 7682 MW; 57F231379AFEEA3C CRC64;
 Query Match 9.9%; Score 8; DB 1; Length 65;
 Best Local Similarity 100.0%; Pred. No. 0.17;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 35 QKEQQLIR 42
 Db 19 QKEQQLIR 26
 RESULT 2
 SUBA_HUMAN
 ID SUBA_HUMAN STANDARD; PRT; 284 AA.
 AC 006520;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 15-JUN-2002 (Rel. 41, Last annotation update)
 DE Alcohol sulfotransferase (EC 2.8.2.2) (Hydroxysteroid
 DE sulfotransferase) (HST) (Dehydroepiandrosterone sulfotransferase)
 DE (DHEA-ST) (ST2) (ST2A3).
 GN SULF2A1 OR STD OR HST.
 OS Homo sapiens (human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 ON NCBI_TaxID=9606;
 RX [1]
 RP SEQUENCE FROM N.A., AND SEQUENCE OF 60-64; 104-119 AND 273-284.
 RC TISSUE-Liver;
 RX MEDLINE=93143674; PubMed=7678732;
 RA Comer K.A., Falany J.L., Falany C.N.,
 RT "Cloning and expression of human liver dehydroepiandrosterone
 RT sulfotransferase.",
 RL Biochem. J. 289:233-240(1993).
 RL [2]
 RP SEQUENCE FROM N.A., AND SEQUENCE OF 80-107 AND 176-198.
 RC TISSUE-Liver;
 RX MEDLINE=92269778; PubMed=1588921;
 RA Ohterness D.M., Wieben E.D., Wood T.C., Watson R.W.G., Madden B.J.,
 RA McCormick D.J., Weinshilboum K.M.,
 RT "Human liver dehydroepiandrosterone sulfotransferase: molecular
 RT cloning and expression of cDNA.",
 RL Mol. Pharmacol. 41:865-872(1992).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE-Adrenal gland;
 RX MEDLINE=96034512; PubMed=7589785;

RA Forbes K.J., Hagen M., Coughtrie M.W.H., Glat H.R., Hume R.,
 RT "Human fetal adrenal hydroxysteroid sulphotransferase: cDNA cloning,
 RT stable expression in V79 cells and functional characterisation of the
 RT expressed enzyme.",
 RL Mol. Cell. Endocrinol. 112:53-60(1995).
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95322029; PubMed=7598806;
 RA Liu-the V., Dufort I., Paquet N., Reimnitz G., Labrie F.,
 RT "Structural characterization and expression of the human
 RT dehydroepiandrosterone sulfotransferase gene.",
 RL DNA Cell Biol. 14:511-518(1995).
 RN [5]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95225980; PubMed=7710689;
 RA Ohterness D.M., Her C., Aksoy S., Kimura S., Wieben E.D.,
 RA Weinshilboum R.M.,
 RT "Human dehydroepiandrosterone sulfotransferase gene: molecular
 RT cloning and structural characterization.",
 RL DNA Cell Biol. 14:331-341(1995).
 RN [6]
 RP SEQUENCE FROM N.A.
 RC TISSUE-Liver;
 RX MEDLINE=92392364; PubMed=1520333;
 RA Kong A.-N.T., Yang L., Ma M., Tao D., Bjornsson T.D.,
 RT "Molecular cloning of the alcohol/hydroxysteroid form (hSta) of
 RT sulfotransferase from human liver.",
 RL Biochem. Biophys. Res. Commun. 187:448-454(1992).
 RN [7]
 RP SEQUENCE FROM N.A.
 RC TISSUE-Liver;
 RA Strausberg R.,
 RT Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: CATALYZES THE SULFATION OF STEROIDS AND BILE ACIDS IN
 CC THE LIVER AND ADRENAL GLANDS.
 CC -1- CATALYTIC ACTIVITY: 3'-phosphoadenylylsulfate + an alcohol =
 CC adenosine 3',5'-bisphosphate + an alkyl sulfate.
 CC -1- SUBUNIT: HOMODIMER.
 CC -1- SUBCELLULAR LOCATION: Cytoplasmic.
 CC -1- TISSUE SPECIFICITY: LIVER, ADRENAL AND AT LOWER LEVEL IN THE
 CC KIDNEY. IS PRESENT IN HUMAN FETUS IN HIGHER LEVEL IN THE ADRENAL
 CC THAN THE LIVER AND THE KIDNEY.
 CC -1- PTM: THE N-TERMINUS IS BLOCKED.
 CC -1- MISCELLANEOUS: ESTROGENS PRESENT IN MATERNAL CIRCULATION IS
 CC PREDOMINANTLY DERIVED FROM FETAL DEHYDROEPIANDROSTERONE SULFATE
 CC WHICH IS HYDROLYZED AND METABOLIZED TO ESTROGENS IN PLACENTA.
 CC -1- SIMILARITY: BELONGS TO THE SULFOTRANSFERASE FAMILY.
 CC
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 CC
 DR EMBL: L20000; AAA35758.1; -
 DR EMBL: X70223; CAA49755.1; -
 DR EMBL: U08024; AAA17748.1; -
 DR EMBL: U08025; AAA17750.1; -
 DR EMBL: X84816; CAA59274.1; -
 DR EMBL: L36196; AAA75491.1; -
 DR EMBL: L36197; AAA75491.1; -
 DR EMBL: L36199; AAA75491.1; JOINED.
 DR EMBL: L36193; AAA75491.1; JOINED.
 DR EMBL: L36194; AAA75491.1; JOINED.
 DR EMBL: L36195; AAA75491.1; JOINED.
 DR EMBL: U13061; AAC51353.1; -
 DR EMBL: U13056; AAC51353.1; JOINED.
 DR EMBL: U13057; AAC51353.1; JOINED.
 DR EMBL: U13058; AAC51353.1; JOINED.
 DR EMBL: U13059; AAC51353.1; JOINED.
 DR EMBL: U13060; AAC51353.1; JOINED.

DR EMBL: S43859; AAB23169.2; -
 DR EMBL: BC020755; AAH20755.1; -
 DR HSSP: P50224; ICM.
 DR Genew; HGNC:11458; SULT2A1.
 DR MIM; 125263; -
 DR InterPro: IPR000863; Sulfotransferase.
 DR Pfam: PF00685; Sulfotransferase; 1.
 DR Prodom: PD001218; Sulfotransferase; 1.
 KM Transferase; Steroid metabolism.
 FT INT_MET 0
 FT BINDING 0
 FT CONFLICT 248 254 PAPS BINDING SITE (POTENTIAL).
 FT CONFLICT 62 62 A -> P (IN REF. 1; AA SEQUENCE).
 FT CONFLICT 89 89 T -> S (IN REF. 1).
 FT CONFLICT 118 118 L -> D (IN REF. 1; AA SEQUENCE).
 FT CONFLICT 158 158 L -> V (IN REF. 6).
 SO SEQUENCE 284 AA; 33648 MW; 3C89C7597833EBA1 CRC64;

Query Match 9.9%; Score 8; DB 1; Length 284;
 Best Local Similarity 100.0%; Pred. No. 0.59;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 48 FRSETLRK 55
 DB 17 FRSETLRK 24

RESULT 3

SUHA_MACFA STANDARD; PRT; 284 AA.
 ID SUHA_MACFA P52842;

DR 01-OCT-1996 (Rel. 34, Created)

DT 01-OCT-1996 (Rel. 34, Last sequence update)

DE 16-OCT-2001 (Rel. 40, Last annotation update)

DE Alcohol sulfotransferase (EC 2.8.2.2) (Hydroxysteroid

GN SULT2A1 OR STD.

OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;

OC Cercopithecoidea; Macaca.

NCBI_TaxID=9541;

RN [1]

SEQUENCE FROM N.A.

RC TISSUE=Liver;

RA Ogura K., Satsukawa M., Kato K., Okuda H., Watabe T.;

Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.

CC -1- FUNCTION: CATALYZES THE SULFATION OF STEROIDS AND BILE ACIDS IN

CC THE LIVER AND ADRENAL GLANDS (BY SIMILARITY).

CC -1- CATALYTIC ACTIVITY: 3'-phosphoadenylylsulfate + an alcohol =

CC adenosine 3',5'-bisphosphate + an alkyl sulfate.

CC -1- SUBUNIT: HOMODIMER (BY SIMILARITY).

CC -1- SIMILARITY: BELONGS TO THE SULFOTRANSFERASE FAMILY.

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CC -----

DR EMBL: D85521; BAA12823.1; -

DR HSSP: P49891; IACU

DR InterPro: IPR000863; Sulfotransferase.

DR Pfam: PF00685; Sulfotransferase; 1.

DR Prodom: PD001218; Sulfotransferase; 1.

DR Transferase; Steroid metabolism.

FT INT_MET 0

FT BINDING 248 254 PAPS BINDING SITE (POTENTIAL).

SO SEQUENCE 284 AA; 33789 MW; 8A8DC56BC0B7A9BD CRC64;

Query Match 9.9%; Score 8; DB 1; Length 284;

Best Local Similarity 100.0%; Pred. No. 0.59;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 48 FRSETLRK 55
 DB 17 FRSETLRK 24

RESULT 4

ALU2_HUMAN STANDARD; PRT; 587 AA.
 ID ALU2_HUMAN P39189;

DR 01-FEB-1995 (Rel. 31, Created)

DT 01-FEB-1995 (Rel. 31, Last sequence update)

DE 16-OCT-2001 (Rel. 40, Last annotation update)

DE Alu subfamily 5B sequence contamination warning entry.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

NCBI_TaxID=9606;

RN [1]

SEQUENCE FROM N.A.

RA MEDLINE=95021758; PubMed=7935834;

RA Claverie J.-M., Makalowski W.;

RT "Alu alert.";

RL Nature 371:752-752(1994).

RN [2]

CONCEPT.

RA MEDLINE=92241891; PubMed=1572661;

RA Claverie J.-M.;

RT "Identifying coding exons by similarity search: alu-derived and other

RL potentially misleading protein sequences.";

RL Genomics 12:838-841(1992).

RN [3]

ALU FAMILIES CLASSIFICATION.

RA MEDLINE=88333009; PubMed=3138422;

RA Quesada Y.;

RT "The Alu family developed through successive waves of fixation

RL closely connected with primate lineage history.";

RL J. Mol. Evol. 27:194-202(1988).

RN [4]

ALU FAMILIES CLASSIFICATION.

RA MEDLINE=91178815; PubMed=1706781;

RA Jurka J., Milosavljevic A.;

RT "Reconstruction and analysis of human Alu genes.";

RL J. Mol. Evol. 32:105-121(1991).

CC -1- MISCELLANEOUS: VARIOUS ANALYSES (SEE REF.3 AND REF.4) INDICATE

CC THAT ALU REPEATS FALL INTO 8 SUBFAMILIES. THEREFORE, 8 ALU WARNING

CC FRAMES CONCEPTUAL TRANSLATIONS OF EACH OF THESE CLASSES OF ALU

CC REPEATS.

CC -1- MISCELLANEOUS: ISOLATED 'X' INDICATES THE PRESENCE OF A STOP

CC CODON. 'XXX' IS USED TO SEPARATE THE VARIOUS TRANSLATION PHASES.

CC -1- CAUTION: THIS ALU ENTRY IS PROVIDED IN ORDER TO AVOID THE FURTHER

CC POLLUTION OF PROTEIN SEQUENCE DATABASES WITH ALU-DERIVED AMINO

CC ACID SEQUENCES.

CC -1- CAUTION: ALU REPETITIVE SEQUENCES ARE INTERSPERSED IN HUMAN AND

CC PRIMATE GENOMES WITH AN AVERAGE SPACING OF 4 KB. SOME OF THEM ARE

CC ACTIVELY TRANSCRIBED BY POL III. NORMAL TRANSCRIPTS MAY CONTAIN

CC ALU-DERIVED SEQUENCES IN 5' OR 3' UNTRANSLATED REGIONS. HOWEVER,

CC CNA LIBRARIES ALSO CONTAIN PARTIAL AND/OR REARRANGED CNAS

CC LIGATED WITH ALU-DERIVED SEQUENCE IN ANY ORIENTATION. ALTHOUGH ALU

CC ELEMENTS (ESPECIALLY SITUATED ON THE COMPLEMENTARY STRAND) HAVE A

CC GREAT POTENTIAL TO CREATE ADDITIONAL/ALTERNATIVE EXONS,

CC CONSIDERATION SHOULD BE GIVEN TO THE POSSIBILITY THAT THE PRESENCE

CC OF AN ALU IN AN OPEN READING FRAME MAY HAVE RESULTED FROM A

CC CLONING ARTIFACT OR MAY BE DUE TO MISINTERPRETATION OF SEQUENCING

CC DATA. THIS POINT HAS BEEN OVERLOOKED ON SEVERAL OCCASIONS, WITH

CC THE CONSEQUENCE OF ERRONEOUS ALU-DERIVED AMINO ACID SEQUENCES

CC BEING REPORTED.

CC -1- CAUTION: ANY SIGNIFICANT SIMILARITY OF A PUTATIVE PROTEIN SEQUENCE

CC WITH AN ALU-TRANSLATED ENTRY MUST BE TAKEN AS A WARNING THAT A

CC PART OF ALU REPEAT MAY HAVE BEEN ARTIFICIALLY INCLUDED IN THE

CC CODING NUCLEOTIDE SEQUENCE.

CC -----

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CC -----

DR EMBL: U14568; -; NOT_ANNOTATED_CDS.

KM Hypothetical protein.

FT DOMAIN 1 96 FRAME-1.

FT DOMAIN 100 194 FRAME-2.

FT DOMAIN 198 292 FRAME-3.

FT DOMAIN 296 391 FRAME-4.

FT DOMAIN 395 489 FRAME-5.

FT DOMAIN 493 587 FRAME-6.

SO SEQUENCE 587 AA; 63703 MW; 3EAB3BE3E3929203 CRC64;

Query Match 9.9%; Score 8; DB 1; Length 587;

Best Local Similarity 100.0%; Pred. No. 1.1;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 SIGDRL 26

DB 183 SIGDRL 190

RESULT 5

ALU3_HUMAN

ID ALU3_HUMAN STANDARD; PRT; 587 AA.

AC P39190;

DT 01-FEB-1995 (Rel. 31, Created)

DT 01-FEB-1995 (Rel. 31, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)

DE Alu subfamily SBI sequence contamination warning entry.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=95021758; PubMed=7935834;

RA Claverie J.-M., Makalowski W.;

RT "Alu alert.";

RL Nature 371:752-752(1994).

RN [2]

RP CONCEPT.

RX MEDLINE=92241891; PubMed=1572661;

RA Claverie J.-M.;

RT "Identifying coding exons by similarity search: alu-derived and other

RT potentially misleading protein sequences.";

RL Genomics 12:838-841(1992).

RN [3]

RP ALU FAMILIES CLASSIFICATION.

RX MEDLINE=88333009; PubMed=3138422;

RA Quentin Y.;

RT "The Alu family developed through successive waves of fixation

RT closely connected with primate lineage history.";

RL J. Mol. Evol. 27:194-202(1988).

RN [4]

RP ALU FAMILIES CLASSIFICATION.

RX MEDLINE=91178815; PubMed=1706781;

RA Jurka J., Milosavljevic A.;

RT "Reconstruction and analysis of human Alu genes.";

RL J. Mol. Evol. 32:105-121(1991).

CC -1- MISCELLANEOUS: VARIOUS ANALYSES (SEE REF. 3 AND REF. 4) INDICATE

CC THAT ALU REPEATS FALL INTO 8 SUBFAMILIES. THEREFORE, 8 ALU WARNING

CC CONSENSUS SEQUENCES HAVE BEEN CONSTITUTED THAT CONTAIN ALL SIX

CC FRAMES CONCEPTUAL TRANSLATIONS OF EACH OF THESE CLASSES OF ALU

CC REPEATS.

CC -1- MISCELLANEOUS: ISOLATED 'X' INDICATES THE PRESENCE OF A STOP

CC CODON, 'xxx' IS USED TO SEPARATE THE VARIOUS TRANSLATION PHASES.

CC -1- CAUTION: THIS ALU ENTRY IS PROVIDED IN ORDER TO AVOID THE FURTHER

CC POLLUTION OF PROTEIN SEQUENCE DATABASES WITH ALU-DERIVED AMINO

CC ACID SEQUENCES.

CC -1- CAUTION: ALU REPETITIVE SEQUENCES ARE INTERSPERSED IN HUMAN AND

CC PRIMATE GENOMES WITH AN AVERAGE SPACING OF 4 KB. SOME OF THEM ARE

CC ACTIVELY TRANSCRIBED BY POL. III. NORMAL TRANSCRIPTS MAY CONTAIN

CC ALU-DERIVED SEQUENCES IN 5' OR 3' UNTRANSLATED REGIONS. HOWEVER,

CC CDNA LIBRARIES ALSO CONTAIN PARTIAL AND/OR REARRANGED CDNAS

CC LIGATED WITH ALU-DERIVED SEQUENCE IN ANY ORIENTATION. ALTHOUGH ALU

CC ELEMENTS (ESPECIALLY SITUATED ON THE COMPLEMENTARY STRAND) HAVE A

CC GREAT POTENTIAL TO CREATE ADDITIONAL/ALTERNATIVE EXONS,

CC CONSIDERATION SHOULD BE GIVEN TO THE POSSIBILITY THAT THE PRESENCE

CC OF AN ALU IN AN OPEN READING FRAME MAY HAVE RESULTED FROM A

CC CLONING ARTIFACT OR MAY BE DUE TO MISINTERPRETATION OF SEQUENCING

CC DATA. THIS POINT HAS BEEN OVERLOOKED ON SEVERAL OCCASIONS. WITH

CC THE CONSEQUENCE OF ERRONEOUS ALU-DERIVED AMINO ACID SEQUENCES

CC BEING REPORTED.

CC -1- CAUTION: ANY SIGNIFICANT SIMILARITY OF A PUTATIVE PROTEIN SEQUENCE

CC WITH AN ALU-TRANSLATED ENTRY MUST BE TAKEN AS A WARNING THAT A

CC PART OF ALU REPEAT MAY HAVE BEEN ARTIFACTUALLY INCLUDED IN THE

CC CODING NUCLEOTIDE SEQUENCE.

CC -----

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CC -----

DR EMBL: U14569; -; NOT_ANNOTATED_CDS.

KM Hypothetical protein.

FT DOMAIN 1 96 FRAME-1.

FT DOMAIN 100 194 FRAME-2.

FT DOMAIN 198 292 FRAME-3.

FT DOMAIN 296 391 FRAME-4.

FT DOMAIN 395 489 FRAME-5.

FT DOMAIN 493 587 FRAME-6.

SO SEQUENCE 587 AA; 63573 MW; 85C4155726DEF235 CRC64;

Query Match 9.9%; Score 8; DB 1; Length 587;

Best Local Similarity 100.0%; Pred. No. 1.1;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 SIGDRL 26

DB 183 SIGDRL 190

RESULT 6

ALU8_HUMAN

ID ALU8_HUMAN STANDARD; PRT; 591 AA.

AC P39195;

DT 01-FEB-1995 (Rel. 31, Created)

DT 01-FEB-1995 (Rel. 31, Last sequence update)

DT 15-JUN-2002 (Rel. 41, Last annotation update)

DE Alu subfamily SX sequence contamination warning entry.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=95021758; PubMed=7935834;

RA Claverie J.-M., Makalowski W.;

RT "Alu alert.";

RL Nature 371:752-752(1994).

RN [2]

RP CONCEPT.

RX MEDLINE=92241891; PubMed=1572661;

RA Claverie J.-M.;

RT "Identifying coding exons by similarity search: alu-derived and other

RT potentially misleading protein sequences.";
 RL Genomics 12:838-841(1992).
 RN [3]
 RP ALU FAMILIES CLASSIFICATION.
 RX MEDLINE=88333009; PubMed=3138422;
 RA Quentin Y.;
 RT "The Alu family developed through successive waves of fixation
 RT closely connected with primate lineage history.";
 RL J. Mol. Evol. 27:194-202(1988).
 RN [4]
 RP ALU FAMILIES CLASSIFICATION.
 RX MEDLINE=91178815; PubMed=1706781;
 RA Jurka J., Milosavljevic A.;
 RT "Reconstruction and analysis of human Alu genes.";
 RL J. Mol. Evol. 32:105-121(1991).
 CC -1- MISCELLANEOUS: VARIOUS ANALYSES (SEE REF.3 AND REF.4) INDICATE
 CC THAT ALU REPEATS FALL INTO 8 SUBFAMILIES. THEREFORE, 8 ALU WARNING
 CC CONSENSUS SEQUENCES HAVE BEEN CONSTITUTED THAT CONTAIN ALL SIX
 CC FRAMES CONCEPTUAL TRANSLATIONS OF EACH OF THESE CLASSES OF ALU
 CC REPEATS.
 CC -1- MISCELLANEOUS: ISOLATED 'X' INDICATES THE PRESENCE OF A STOP
 CC CODON, 'XXX' IS USED TO SEPARATE THE VARIOUS TRANSLATION PHASES.
 CC -1- CAUTION: THIS ALU ENTRY IS PROVIDED IN ORDER TO AVOID THE FURTHER
 CC POLLUTION OF PROTEIN SEQUENCE DATABASES WITH ALU-DERIVED AMINO
 CC ACID SEQUENCES.
 CC -1- CAUTION: ALU REPETITIVE SEQUENCES ARE INTERSPERSED IN HUMAN AND
 CC PRIMATE GENOMES WITH AN AVERAGE SPACING OF 4 KB. SOME OF THEM ARE
 CC ACTIVELY TRANSCRIBED BY POL III. NORMAL TRANSCRIPTS MAY CONTAIN
 CC ALU-DERIVED SEQUENCES IN 5' OR 3' UNTRANSLATED REGIONS. HOWEVER,
 CC CDNA LIBRARIES ALSO CONTAIN PARTIAL AND/OR REARRANGED CDNAS
 CC LIGATED WITH ALU-DERIVED SEQUENCE IN ANY ORIENTATION. ALTHOUGH ALU
 CC ELEMENTS (ESPECIALLY SITUATED ON THE COMPLEMENTARY STRAND) HAVE A
 CC GREAT POTENTIAL TO CREATE ADDITIONAL/ALTERNATIVE EXONS,
 CC CONSIDERATION SHOULD BE GIVEN TO THE POSSIBILITY THAT THE PRESENCE
 CC OF AN ALU IN AN OPEN READING FRAME MAY HAVE RESULTED FROM A
 CC CLONING ARTIFACT OR MAY BE DUE TO MISINTERPRETATION OF SEQUENCING
 CC DATA. THIS POINT HAS BEEN OVERLOOKED ON SEVERAL OCCASIONS, WITH
 CC THE CONSEQUENCE OF ERRONEOUS ALU-DERIVED AMINO ACID SEQUENCES
 CC BEING REPORTED.
 CC -1- CAUTION: ANY SIGNIFICANT SIMILARITY OF A PUTATIVE PROTEIN SEQUENCE
 CC WITH AN ALU-TRANSLATED ENTRY MUST BE TAKEN AS A WARNING THAT A
 CC PART OF ALU REPEAT MAY HAVE BEEN ARTIFACTUALLY INCLUDED IN THE
 CC CODING NUCLEOTIDE SEQUENCE.
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: U14574; ; NOT_ANNOTATED_CDS.
 KW Hypothetical protein.
 FT DOMAIN 1 96 FRAME-1.
 FT DOMAIN 100 195 FRAME-2.
 FT DOMAIN 199 294 FRAME-3.
 FT DOMAIN 298 393 FRAME-4.
 FT DOMAIN 397 492 FRAME-5.
 FT DOMAIN 496 591 FRAME-6.
 SQ SEQUENCE 591 AA; 64395 MW; AC8154AD8A6B280 CRC64;
 Query Match 9.9%; Score 8; DB 1; Length 591;
 Best Local Similarity 100.0%; Pred. No. 1.1;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 7
 FLHD_XENNE

ID FLHD_XENNE STANDARD; PRT; 116 AA.
 AC G9X9F2;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 15-JUN-2002 (Rel. 41, Last annotation update)
 DE Flagellar transcriptional activator flhd.
 GN FLHD.
 OS Xenorhabdus nematophilus.
 OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 OC Xenorhabdus.
 OX NCBI_TaxID=628;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=FL;
 RA Givaudan A.G., Janois A.;
 RT "flhnc gene disruptions leads to pleiotropic phenotypes.";
 RL Submitted (NOV-1998) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: Transcriptional activator. Together with flnc it acts as
 CC a compound sigma factor that activates class 2 flagellar genes (By
 CC similarity).
 CC -1- SIMILARITY: BELONGS TO THE FLHD FAMILY.
 CC -----
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 CC -----
 CC EMBL: AJ012828; CAB41407.1; ;
 KW DNA-directed RNA polymerase; DNA-binding; Flagella.
 SQ SEQUENCE 116 AA; 13303 MW; 89DBF8175532828 CRC64;
 Query Match 8.6%; Score 7; DB 1; Length 116;
 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 8
 INVO_CANFA STANDARD; PRT; 285 AA.
 AC P18174;
 DT 01-NOV-1990 (Rel. 16, Created)
 DT 01-NOV-1990 (Rel. 16, Last sequence update)
 DT 15-JUN-2002 (Rel. 41, Last annotation update)
 DE Involucrin.
 GN IVL.
 OS Canis familiaris (Dog).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
 OX NCBI_TaxID=9615;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90348475; PubMed=2385171;
 RA Tseeng H., Green H.;
 RT "The involucrin genes of pig and dog: comparison of their segments of
 RT repeats with those of primates and higher primates.";
 RL Mol. Biol. Evol. 7:293-302(1990).
 CC -1- FUNCTION: Part of the insoluble cornified cell envelope (CE) of
 CC stratified squamous epithelia.
 CC -1- SUBCELLULAR LOCATION: Cytoplasmic. Constituent of the scaffolding
 CC of the cornified envelope.
 CC -1- TISSUE SPECIFICITY: Keratinocytes of epidermis and other
 CC stratified squamous epithelia.
 CC -1- PTM: Substrate of transglutaminase. Specific glutamines or lysines
 CC are cross-linked to keratins, desmoplakin and to inter involucrin
 CC molecules.

CC -1- SIMILARITY: BELONGS TO THE INVOLUCRIN FAMILY.
 CC -----
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 CC -----
 DR EMBL: M34442; AAA30853.1; -
 DR InterPro: IPR002360; Involucrin.
 DR PROSITE: PS00795; INVOLUCRIN; 1.
 SW KeraLinization; Repeat.
 SQ SEQUENCE 285 AA; 33384 MW; DCE1BD88B9248BEA CRC64;
 Query Match 8.6%; Score 7; DB 1; Length 285;
 Best Local Similarity 100.0%; Pred. No. 6.5;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 32 KQOQKQEQ 38
 Db 198 KQOQKQEQ 204
 RESULT 9
 Y687_HAEIN STANDARD; PRT; 304 AA.
 ID Y687_HAEIN
 AC P71356;
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-JUN-2002 (Rel. 41, Last annotation update)
 DE Hypothetical transport protein H10687.
 GN H10687.
 OS Haemophilus influenzae.
 OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
 CC Haemophilus.
 CC NBL_TaxID=727;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-Rd / KW20 / ATCC 51907;
 RX MEDLINE=95350630; PubMed=7542800;
 RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
 RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
 RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
 RA Scott J.D., Shiley R., Liu L.-I., Glodek A., Kelley J.M.,
 RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
 RA Uterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
 RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
 RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
 RA Venter J.C.;
 RT Whole-genome random sequencing and assembly of Haemophilus influenzae
 RT Rd.";
 RL Science 269:496-512(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (Probable).
 CC -1- SIMILARITY: BELONGS TO THE EAMA TRANSPORTER FAMILY.
 CC -----
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 CC -----
 DR EMBL: U32751; AAC2347.1; -
 DR TIGR: H10687; -
 DR InterPro: IPR000620; DUF6.
 DR Pfam: PF00892; DUF6; 2.
 KW Hypothetical protein; Transport; Transmembrane; Complete proteome.
 FT TRANSMEM 9 29 POTENTIAL.
 FT TRANSMEM 67 87 POTENTIAL.
 FT TRANSMEM 100 120 POTENTIAL.

FT TRANSMEM 131 151 POTENTIAL.
 FT TRANSMEM 159 179 POTENTIAL.
 FT TRANSMEM 189 209 POTENTIAL.
 FT TRANSMEM 222 242 POTENTIAL.
 FT TRANSMEM 252 272 POTENTIAL.
 FT TRANSMEM 278 298 POTENTIAL.
 SQ SEQUENCE 304 AA; 33887 MW; CC7095529EB8E4FB3 CRC64;
 Query Match 8.6%; Score 7; DB 1; Length 304;
 Best Local Similarity 100.0%; Pred. No. 6.9;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 7 LFITYLFI 13
 Db 71 LFITYLFI 77
 RESULT 10
 YK68_YEAST STANDARD; PRT; 337 AA.
 ID YK68_YEAST
 AC P36164;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DE Hypothetical 38.3 kDa protein in PRP16-SRP40 intergenic region.
 GN YK088C OR YK0408.
 OS Saccharomyces cerevisiae (Baker's yeast).
 CC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 CC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 CC NBL_TaxID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=94262327; PubMed=8203164;
 RA Garcia-Cantalejo J., Baladron V., Esteban P.F., Santos M.A., Bou G.,
 RA Remacha M.A., Revuelta J.L., Ballesta J.P.G., Jimenez A., del Rey F.;
 RT "The complete sequence of an 18,002 bp segment of Saccharomyces
 RT cerevisiae chromosome XI contains the HBS1, MRP-L20 and PRP16 genes,
 RT and six new open reading frames.";
 RL Yeast 10:231-245(1994).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
 CC -----
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 CC -----
 DR EMBL: Z27116; CAA81639.1; -
 DR EMBL: Z28313; CAA82167.1; -
 DR PIR: S38166; S38166.
 DR PIR: S39129; S39129.
 DR SGD: S0001796; YKR088C.
 KW Hypothetical protein; Transmembrane.
 FT TRANSMEM 96 115 POTENTIAL.
 FT TRANSMEM 138 162 POTENTIAL.
 FT TRANSMEM 173 191 POTENTIAL.
 FT TRANSMEM 222 246 POTENTIAL.
 FT TRANSMEM 253 271 POTENTIAL.
 FT TRANSMEM 287 309 POTENTIAL.
 SQ SEQUENCE 337 AA; 38311 MW; 7EA95DD4E5A77FE CRC64;
 Query Match 8.6%; Score 7; DB 1; Length 337;
 Best Local Similarity 100.0%; Pred. No. 7.5;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 7 LFITYLFI 13
 Db 264 LFITYLFI 270
 RESULT 11

CITC_ECOLI
ID CITC_ECOLI STANDARD; PRT; 352 AA.
AC P77390; 054337; 09R2T4;
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE [citr-35]-lyase [citr-35]-lyase (EC 6.2.1.22) (citr-35-lyase synthetase) (Acetate:SH-citr-35-lyase synthetase).
OS Citric acid synthetase.
GN Citric acid synthetase.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
OX NCBI_Taxid=562;
[1]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RA MEDLINE=97426617; PubMed=9278503;
RA Blatter F.R., Plunkett G., III, Bloch C.A., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12.";
RL Science 277:1453-1474(1997).
[2]
RP SEQUENCE FROM N.A.
RA Chung E., Allen E., Araujo R., Aparicio A., Davis K., Duncan M.,
RA Federapoli N., Hyman R., Kalman S., Komp C., Kurdi O., Lew H., Lin D.,
RA Nemat A., Oefner P., Roberts D., Schramm S., Davis R.W.;
RL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.
[3]
RP SEQUENCE FROM N.A.
RC STRAIN=K12;
RA MEDLINE=97061202; PubMed=8905232;
RA Oshima T., Alba H., Baba T., Fujita K., Hayashi K., Honjo A.,
RA Ikeno K., Inada T., Itoh T., Kajihara M., Kanai K., Kashimoto K.,
RA Kimura S., Kitagawa M., Makino K., Masuda S., Miki T., Mizobuchi K.,
RA Mori H., Motomura K., Nakamura Y., Nishimoto H., Nishio Y., Saito N.,
RA Sempel G., Seki Y., Tagami H., Takemoto K., Wada C., Yamamoto Y.,
RA Yano M., Horiuchi T.;
RT "A 718-kb DNA sequence of the Escherichia coli K-12 genome
corresponding to the 12.7-28.0 min region on the linkage map.";
RL DNA Res. 3:137-155(1996).
[4]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RA Ingmer H., Cohen S.N.;
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: ACETYLATION OF PROSTHETIC GROUP (2-(5'-PHOSPHORIBOSYL)-
3-DEPHOSPHOCOENZYME-A) OF THE GAMMA SUBUNIT OF CITRATE LYASE.
CC -1- CATALYTIC ACTIVITY: ATP + acetate + [citr-35]-lyase
(thiol form) = AMP + diphosphate + [citr-35]-lyase
(acetyl form).
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CC -----
DR EMBL; AEO00166; AAC73719.1; ALT_INIT.
DR EMBL; U82598; AAB40818.1; ALT_INIT.
DR EMBL; D90702; BAA35254.1; ALT_INIT.
DR EMBL; D90703; BAA35260.1; ALT_INIT.
DR EMBL; U46667; AAC28950.1; --
DR EcoGene; EGI3645; citC.
DR InterPro; IPR005216; cit_ly_11gase.
DR InterPro; IPR004821; cit_ly_11gase.
DR InterPro; IPR000183; GCSacetyltransf.
DR Pfam; PF00583; Acetyltransf. 1.
DR TIGRfams; TIGR00124; cit_ly_11gase. 1.
DR TIGRfams; TIGR00125; cit_ly_11gase. 1.

KW Ligase: Complete proteome.
SQ SEQUENCE 352 AA; 40077 MW; F5894FCD0F06518 CRC64;
Query Match 8.6%; Score 7; DB 1; Length 352;
Best Local Similarity 100.0%; Pred. No. 7.8;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 4 STHELIT 10
|||||
Db 90 STHELIT 96
RESULT 12
ID ROX1_YEAST STANDARD; PRT; 368 AA.
AC P25042;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE ROX1 repressor (Hypoxic function repressor) (Heme-dependent repression factor).
GN ROX1 OR YPR065W OR YP9499.20.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
OX NCBI_Taxid=4932;
[1]
RP SEQUENCE FROM N.A.
RC MEDLINE=94019282; PubMed=8413209;
RA Balasubramanian B., Lowry C.V., Zlotner R.S.;
RT "The RoX1 repressor of the Saccharomyces cerevisiae hypoxic genes is
a specific DNA-binding protein with a high-mobility-group motif.";
RL Mol. Cell. Biol. 13:6071-6076(1993).
[2]
RP SEQUENCE FROM N.A.
RC STRAIN=S288C / AB972;
RA MEDLINE=97313271; PubMed=9169875;
RA Bussey H., Storms R.K., Ahmed A., Albermann K., Allen E., Ansgore W.,
RA Araujo R., Aparicio A., Barrell B.G., Badcock K., Benes V.,
RA Botstein D., Bowman S., Bruckner M., Carpenter J., Cherry J.M.,
RA Chung E., Churcher C.M., Coster F., Davis K., Davis R.W.,
RA Dietrich F.S., Delius H., DiPaolo T., Dubois E., Duesterhoeft A.,
RA Duncan M., Floeth M., Fortin N., Friesen J.D., Fritz C., Goffeau A.,
RA Hall J., Hebling U., Heumann K., Hilbert H., Hillier L.,
RA Hunnicke-Smith S., Hyman R., Johnston M., Kalman S., Kleine K.,
RA Komp C., Kurdi O., Lashkari D., Lew H., Lin A., Lin D., Louis E.J.,
RA Marathe R., Messenguy F., Mewes H.-W., Mittlepelt S., Mostl D.,
RA Mueller-Auer S., Namath A., Nentwich U., Oefner P., Pearson D.,
RA Petel F.X., Pohl T.M., Purnelle D., Schafer M., Scharfe M.,
RA Scherens B., Schramm S., Schroeder M., Sidou A.M., Tietelin H.,
RA Urrestegui L.A., Ushinsky S., Vierendeels F., Vissers S., Voss H.,
RA Walsh S.V., Wandut R., Wang Y., Wedler E., Wedler H., Winnett E.,
RA Zhong W.W., Zollner A., Vo D.H., Hani J.;
RT "The nucleotide sequence of Saccharomyces cerevisiae chromosome XVI.";
RL Nature 387:103-105(1997).
[3]
RP CHARACTERIZATION.
RC MEDLINE=96174644; PubMed=8600445;
RA di Flumeri C., Liston P., Acheson N.H., Keng T.;
RT "The HMG domain of the RoX1 protein mediates repression of HEM13
through overlapping DNA binding and oligomerization functions.";
RL Nucleic Acids Res. 24:808-815(1996).
CC -1- FUNCTION: TRANSCRIPTION FACTOR THAT REPRESENTS THE EXPRESSION OF
HEM13, COX5B, ANH1, CYC7 OR AAC3 (HYPOXIC FUNCTION). BINDS TO THE
DNA SEQUENCE 5'-RRRTACAGAC-3'.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- SIMILARITY: CONTAINS 1 HMG BOX.
CC -1- INDUCTION: BY HEME.
CC -----
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DR EMBL: X60458; CAA42991.1; -;
 DR EMBL: Z49219; CAA89182.1; -;
 DR EMBL: Z71255; CAA94973.1; -;
 DR PIR: S17015; S17015.
 DR HSSP: Q05066; 1HR.
 DR TRANSFAC: T01286; -;
 DR SGD: S0006269; ROX1.
 DR Interpro: IPR000910; HMG_12_box.
 DR Pfam: PF00505; HMG_box; 1.
 DR SMART: SM00398; HMG; 1.
 KW Transcription regulation; Repressor; DNA-binding; Nuclear protein.
 FT DNA_BIND 14 83 HMG BOX.
 FT DOMAIN 102 123 GLN-RICH.
 SQ SEQUENCE 368 AA; 41838 MW; 3B27442D7DEE3DBD CRC64;

Query Match 8.6%; Score 7; DB 1; Length 368;
 Best Local Similarity 100.0%; Pred. No. 8.1;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 33 QOOKEOO 39
 Db 106 QOOKEOO 112

RESULT 13
 ALU1_HUMAN STANDARD; PRT; 591 AA.
 AC P39186;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Alu subfamily J sequence contamination warning entry.
 OS Homo sapiens (human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95021758; PubMed=7935834;
 RA Claverie J.-M., Makalowski W.;
 RL "Alu alert.";
 RL Nature 371:752-752(1994).
 RN [2]
 RP CONCEPT.
 RX MEDLINE=92241891; PubMed=1572661;
 RA Claverie J.-M.;
 RT "Identifying coding exons by similarity search: alu-derived and other
 RT potentially misleading protein sequences.";
 RL Genomics 12:838-841(1992).
 RN [3]
 RP ALU FAMILIES CLASSIFICATION.
 RX MEDLINE=88333009; PubMed=3138422;
 RA Quentin Y.;
 RT "The Alu family developed through successive waves of fixation
 RT closely connected with primate lineage history.";
 RL J. Mol. Evol. 27:194-202(1988).
 RN [4]
 RP ALU FAMILIES CLASSIFICATION.
 RX MEDLINE=91178815; PubMed=1706781;
 RA Jurka J., Milosavljevic A.;
 RT "Reconstruction and analysis of human Alu genes.";
 RL J. Mol. Evol. 32:105-121(1991).
 CC -1- MISCELLANEOUS: VARIOUS ANALYSES (SEE REF. 3 AND REF. 4) INDICATE
 CC THAT ALU REPEATS FALL INTO 8 SUBFAMILIES. THEREFORE, 8 ALU WARNING
 CC CONSENSUS SEQUENCES HAVE BEEN CONSTITUTED THAT CONTAIN ALL SIX
 CC FRAMES CONCEPTUAL TRANSLATIONS OF EACH OF THESE CLASSES OF ALU
 CC REPEATS.
 CC -1- MISCELLANEOUS: ISOLATED 'X' INDICATES THE PRESENCE OF A STOP
 CC CODON, 'xxx' IS USED TO SEPARATE THE VARIOUS TRANSLATION PHASES.

CC -1- CAUTION: THIS ALU ENTRY IS PROVIDED IN ORDER TO AVOID THE FURTHER
 CC POLLUTION OF PROTEIN SEQUENCE DATABASES WITH ALU-DERIVED AMINO
 CC ACID SEQUENCES.

CC -1- CAUTION: ALU REPETITIVE SEQUENCES ARE INTERSPERSED IN HUMAN AND
 CC PRIMATE GENOMES WITH AN AVERAGE SPACING OF 4 KB. SOME OF THEM ARE
 CC ACTIVELY TRANSCRIBED BY POL. III. NORMAL TRANSCRIPTS MAY CONTAIN
 CC ALU-DERIVED SEQUENCES IN 5' OR 3' UNTRANSLATED REGIONS. HOWEVER,
 CC CDNA LIBRARIES ALSO CONTAIN PARTIAL AND/OR REARRANGED CDNAS
 CC LIGATED WITH ALU-DERIVED SEQUENCE IN ANY ORIENTATION. ALTHOUGH ALU
 CC ELEMENTS (ESPECIALLY SITUATED ON THE COMPLEMENTARY STRAND) HAVE A
 CC GREAT POTENTIAL TO CREATE ADDITIONAL/ALTERNATIVE EXONS.
 CC CONSIDERATION SHOULD BE GIVEN TO THE POSSIBILITY THAT THE PRESENCE
 CC OF AN ALU IN AN OPEN READING FRAME MAY HAVE RESULTED FROM A
 CC CLONING ARTIFACT OR MAY BE DUE TO MISINTERPRETATION OF SEQUENCING
 CC DATA. THIS POINT HAS BEEN OVERLOOKED ON SEVERAL OCCASIONS, WITH
 CC THE CONSEQUENCE OF ERRONEOUS ALU-DERIVED AMINO ACID SEQUENCES
 CC BEING REPORTED.

CC -1- CAUTION: ANY SIGNIFICANT SIMILARITY OF A PUTATIVE PROTEIN SEQUENCE
 CC WITH AN ALU-TRANSLATED ENTRY MUST BE TAKEN AS A WARNING THAT A
 CC PART OF ALU REPEAT MAY HAVE BEEN ARTIFICIALLY INCLUDED IN THE
 CC CODING NUCLEOTIDE SEQUENCE.

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CC EMBL: U14567; -; NOT_ANNOTATED_CDS.
 KW Hypothetical protein.
 DR DOMAIN 1 96 FRAME-1.
 FT DOMAIN 100 195 FRAME-2.
 FT DOMAIN 199 294 FRAME-3.
 FT DOMAIN 298 393 FRAME-4.
 FT DOMAIN 397 492 FRAME-5.
 FT DOMAIN 496 591 FRAME-6.
 SQ SEQUENCE 591 AA; 655D395735519D95 CRC64;

Query Match 8.6%; Score 7; DB 1; Length 591;
 Best Local Similarity 100.0%; Pred. No. 12;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 19 SLGDRAR 25
 Db 85 SLGDRAR 91

RESULT 14
 ALU4_HUMAN STANDARD; PRT; 603 AA.
 AC P39191;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Alu subfamily SB2 sequence contamination warning entry.
 OS Homo sapiens (human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95021758; PubMed=7935834;
 RA Claverie J.-M., Makalowski W.;
 RL "Alu alert.";
 RL Nature 371:752-752(1994).
 RN [2]
 RP CONCEPT.
 RX MEDLINE=92241891; PubMed=1572661;
 RA Claverie J.-M.;
 RT "Identifying coding exons by similarity search: alu-derived and other
 RT potentially misleading protein sequences.";

RL Genomics 12:838-841(1992).
 RN [3]
 RP ALU FAMILIES CLASSIFICATION.
 RX MEDLINE-88333009; PubMed-3138422;
 RA Quentlin Y.;
 RT "The Alu family developed through successive waves of fixation
 closely connected with primate lineage history.";
 RL J. Mol. Evol. 27:194-202(1988).
 RN [4]
 RP ALU FAMILIES CLASSIFICATION.
 RX MEDLINE-91178815; PubMed-1706781;
 RA Jurka J., Miosavljevic A.;
 RT "Reconstruction and analysis of human Alu genes";
 RL J. Mol. Evol. 32:105-121(1991).
 CC -1- MISCELLANEOUS: VARIOUS ANALYSES (SEE REF.3 AND REF.4) INDICATE
 THAT ALU REPEATS FALL INTO 8 SUBFAMILIES. THEREFORE, 8 ALU WARNING
 CC CONSENSUS SEQUENCES HAVE BEEN CONSTITUTED THAT CONTAIN ALL SIX
 CC FRAMES CONCEPTUAL TRANSLATIONS OF EACH OF THESE CLASSES OF ALU
 CC REPEATS.
 CC -1- MISCELLANEOUS: ISOLATED 'X' INDICATES THE PRESENCE OF A STOP
 CC CODON, 'XXX' IS USED TO SEPARATE THE VARIOUS TRANSLATION PHASES.
 CC -1- CAUTION: THIS ALU ENTRY IS PROVIDED IN ORDER TO AVOID THE FURTHER
 CC POLLUTION OF PROTEIN SEQUENCE DATABASES WITH ALU-DERIVED AMINO
 CC ACID SEQUENCES.
 CC -1- CAUTION: ALU REPETITIVE SEQUENCES ARE INTERSPERSED IN HUMAN AND
 CC PRIMATE GENOMES WITH AN AVERAGE SPACING OF 4 KB. SOME OF THEM ARE
 CC ACTIVELY TRANSCRIBED BY POL. III. NORMAL TRANSCRIPTS MAY CONTAIN
 CC ALU-DERIVED SEQUENCES IN 5' OR 3' UNTRANSLATED REGIONS. HOWEVER,
 CC CDNA LIBRARIES ALSO CONTAIN PARTIAL AND/OR REARRANGED CDNAS
 CC LIGATED WITH ALU-DERIVED SEQUENCE IN ANY ORIENTATION. ALTHOUGH ALU
 CC ELEMENTS (ESPECIALLY SITUATED ON THE COMPLEMENTARY STRAND) HAVE A
 CC GREAT POTENTIAL TO CREATE ADDITIONAL/ALTERNATIVE EXONS,
 CC CONSIDERATION SHOULD BE GIVEN TO THE POSSIBILITY THAT THE PRESENCE
 CC OF AN ALU IN AN OPEN READING FRAME MAY HAVE RESULTED FROM A
 CC CLONING ARTIFACT OR MAY BE DUE TO MISINTERPRETATION OF SEQUENCING
 CC DATA. THIS POINT HAS BEEN OVERLOOKED ON SEVERAL OCCASIONS, WITH
 CC THE CONSEQUENCE OF ERRONEOUS ALU-DERIVED AMINO ACID SEQUENCES
 CC BEING REPORTED.
 CC -1- CAUTION: ANY SIGNIFICANT SIMILARITY OF A PUTATIVE PROTEIN SEQUENCE
 CC WITH AN ALU-TRANSLATED ENTRY MUST BE TAKEN AS A WARNING THAT A
 CC PART OF AN ALU REPEAT MAY HAVE BEEN ARTIFICIALLY INCLUDED IN THE
 CC CODING NUCLEOTIDE SEQUENCE.
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 CC -----
 CC EMBL; U14570; -; NOT ANNOTATED_CDS.
 DR Hypothetical protein.
 KM DOMAIN 1 98 FRAME-1.
 FT DOMAIN 102 199 FRAME-2.
 FT DOMAIN 202 300 FRAME-3.
 FT DOMAIN 304 401 FRAME-4.
 FT DOMAIN 405 502 FRAME-5.
 FT DOMAIN 506 603 FRAME-6.
 SO SEQUENCE 603 AA; 65272 MW; B8AD0D46BEA114 CRC64;
 Query Match 8.6%; Score 7; DB 1; Length 603;
 Best Local Similarity 100.0%; Pred. No. 12;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 20 LGRARL 26
 DB 88 LGRARL 94
 RESULT 15
 SYL_BACSU
 ID SYL_BACSU STANDARD; PRT; 804 AA.

AC P36430; O34465;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 15-JUN-2002 (Rel. 41, Last annotation update)
 DE Leuyl-tRNA synthetase (EC 6.1.1.4) (Leucine--tRNA ligase) (LeuRS).
 GN LEU5.
 OS Bacillus subtilis.
 CC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
 OX NCBI_TaxID=1423;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE-92283747; PubMed-1317842;
 RA Vander Horn P.B., Zahler S.A.;
 RT "Cloning and nucleotide sequence of the leuyl-tRNA synthetase gene
 of Bacillus subtilis";
 RL J. Bacteriol. 174:3928-3935(1992).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=168;
 RX MEDLINE-98048467; PubMed-9387221;
 RA Lapidus A., Galleron N., Sorokin A., Ehrlich S.D.;
 RT "Sequencing and functional annotation of the Bacillus subtilis genes
 in the 200 kb rnb-dnaB region";
 RL Microbiology 143:3431-3441(1997).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=168;
 RX MEDLINE-98044033; PubMed-9384377;
 RA Kunst F., Ogatawara N., Moszer I., Albertini A.M., Alloni G.,
 RA Azevedo V., Bertolo M.G., Bessieres P., Bolotin A., Borchert S.,
 RA Borriest R., Boutsier L., Brans A., Braun M., Brignell S.C., Bron S.,
 RA Brouillet S., Bruschi C.V., Caldwell B., Capuano V., Carter N.M.,
 RA Choi S.K., Codani J.J., Connerton I.F., Cummings N.J., Daniel R.A.,
 RA Deizel F., Devine K.M., Dusterhoft A., Ehrlich S.D., Emmerson P.T.,
 RA Enlian K.D., Errington J., Fadret C., Ferrari E., Foulger D.,
 RA Fritz C., Fujita M., Fujita Y., Fuma S., Gallizi A., Galleron N.,
 RA Gilm S.Y., Glaser P., Goffeau A., Goldlighty E.J., Grandi G.,
 RA Giuseppe G., Guy B.J., Haga K., Halech J., Harwood C.R., Henaut A.,
 RA Gilbert H., Holsappel S., Hosono S., Hulio M.F., Itaya M., Jones L.,
 RA Joris B., Karamata D., Kasahara Y., Klerr-Blanchard M., Klein C.,
 RA Kodayashi Y., Koetter P., Koningsstein G., Krogh S., Kumano M.,
 RA Kunita K., Lapidus A., Lardinois S., Lauber J., Lazarevic V.,
 RA Lee S.M., Levine A., Liu H., Masuda S., Manuel C., Medigue C.,
 RA Medina N., Mellado R.P., Mizuno M., Mostl D., Nakai S., Noback M.,
 RA Noone D., O'Reilly M., Ogawa K., Ogiwara A., Oudaga B., Park S.H.,
 RA Parro V., Pohl T.M., Portelle D., Portwollik S., Prescott A.M.,
 RA Presacan E., Pujic P., Purnelle B., Rapoport G., Rey M., Reynolds S.,
 RA Rieger M., Rivolta C., Rocha E., Roche B., Rose M., Sadale Y.,
 RA Sato T., Scamlan E., Schleich S., Schroeter R., Scofield F.,
 RA Sekiguchi J., Sekowska A., Seror S.J., Seror P., Shin B.S., Soldo B.,
 RA Sorokin A., Tacconi E., Takagi T., Takahashi H., Takemaru K.,
 RA Takeuchi M., Tamakoshi A., Tanaka T., Terpeira P., Tognoni A.,
 RA Trosato V., Uchiyama S., Vandenbol M., Vannier F., Vassaretti A.,
 RA Viari A., Wandt R., Wedler E., Wedler H., Weitzenecker T.,
 RA Winters P., Wipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,
 RA Yoshida K., Yoshikawa H.F., Zumbstein E., Yoshikawa K., Danchin A.;
 RT "The complete genome sequence of the Gram-positive bacterium Bacillus
 subtilis";
 RL Nature 390:249-256(1997).
 CC -1- CATALYTIC ACTIVITY: ATP + L-leucine + tRNA(Leu) -> AMP +
 CC diphosphate + L-leucyl-tRNA(Leu).
 CC -1- SUBCELLULAR LOCATION: cytoplasmic.
 CC -1- SIMILARITY: BELONGS TO CLASS-I AMINOACYL-tRNA SYNTHETASE FAMILY.
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 CC -----
 CC EMBL; M88581; AAA2571.1; -.

DR EMBL; AF008220; AAC00259.1; -
 DR EMBL; Z99119; CAB15010.1; -
 DR PIR; A41882; A41882.
 DR Subtilist; BG10676; leus.
 DR InterPro; IPR002302; leu-TRNAsyntla.
 DR InterPro; IPR002300; tRNA-synt_1a.
 DR InterPro; IPR001412; tRNA-synt_1.
 DR Pfam; PF00133; tRNA-synt_1; 1.
 DR PRINTS; PR00985; TRNASYNTHLEU.
 DR TIGRFAMs; TIGR00396; leus.bact; 1.
 DR PROSITE; PS00178; AA-TRNA_LIGASE_I; 1.
 DR Aminoacyl-tRNA synthetase; Protein biosynthesis; Ligase; ATP-binding;
 KW Complete proteome.
 FT SITE 40 51 "HIGH" REGION.
 FT SITE 576 580 "KMSKS" REGION.
 FT BINDING 579 579 ATP (BY SIMILARITY).
 FT CONFLICT 186 186 P -> L (IN REF. 1).
 FT CONFLICT 195 195 T -> N (IN REF. 1).
 FT CONFLICT 247 281 RPDTLEGATYVLAPEHALVENTTTAEQKEAVEAY -> DQ
 IRLALHTLSLPRTTHWMTSORSKRKLKLI (IN
 REF. 1).
 SO SEQUENCE 804 AA; 91542 MW; 306FD5A98FE5C47E CRC64;

Query Match 8.68; Score 7; DB 1; Length 804;
 Best Local Similarity 100.08; Pred. No. 16;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 56 TGKKGRR 62
 |||||
 Db 479 TGKKGRR 485

Search completed: May 29, 2003, 15:31:09
 Job time : 23 secs